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NEWS 8 FEB 16 INSPEC Adding Its Own IPC codes and Author's E-mail
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NEWS 9 APR 02 CAS Registry Number Crossover Limits Increased to
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NEWS EXPRESS FEBRUARY 15 10 CURRENT WINDOWS VERSION IS V8.4.2,
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Updated Search

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FULL ESTIMATED COST	0.22	0.22

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STRUCTURE FILE UPDATES: 3 JUN 2010 HIGHEST RN 1226953-63-4
DICTIONARY FILE UPDATES: 3 JUN 2010 HIGHEST RN 1226953-63-4

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=>
Uploading C:\Documents and Settings\brobinson1\My Documents\e-Red Folder\arararararar.str

L1 STRUCTURE UPLOADED

=> s l1
SAMPLE SEARCH INITIATED 16:25:40 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 5685 TO ITERATE

35.2% PROCESSED	2000 ITERATIONS	0 ANSWERS
INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)		
SEARCH TIME: 00.00.01		

FULL FILE PROJECTIONS:	ONLINE	**COMPLETE**
	BATCH	**COMPLETE**
PROJECTED ITERATIONS:	109179 TO	118221
PROJECTED ANSWERS:	0 TO	0

Updated Search

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L2 0 SEA SSS SAM L1

=> s l1 full
THE ESTIMATED SEARCH COST FOR FILE 'REGISTRY' IS 191.05 U.S. DOLLARS
DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N or END:y
FULL SEARCH INITIATED 16:25:46 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 114659 TO ITERATE

100.0% PROCESSED 114659 ITERATIONS 0 ANSWERS
SEARCH TIME: 00.00.07

L3 0 SEA SSS FUL L1

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L4 STRUCTURE UPLOADED

=> s l4
SAMPLE SEARCH INITIATED 16:27:26 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 91 TO ITERATE

100.0% PROCESSED 91 ITERATIONS 0 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 1248 TO 2392
PROJECTED ANSWERS: 0 TO 0

L5 0 SEA SSS SAM L4

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100.0% PROCESSED 1807 ITERATIONS 2 ANSWERS
SEARCH TIME: 00.00.01

L6 2 SEA SSS FUL L4

=> file hcaplus
COST IN U.S. DOLLARS SINCE FILE TOTAL
ENTRY SESSION
FULL ESTIMATED COST 386.02 386.24

FILE 'HCAPLUS' ENTERED AT 16:27:33 ON 05 JUN 2010
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Updated Search

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FILE COVERS 1907 - 5 Jun 2010 VOL 152 ISS 24
FILE LAST UPDATED: 4 Jun 2010 (20100604/ED)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Apr 2010
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Apr 2010

HCAplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2010.

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This file contains CAS Registry Numbers for easy and accurate substance identification.

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L7          3 L6

=> s l7 and lightner, j?/au
          55 LIGHTNER, J?/AU
L8          0 L7 AND LIGHTNER, J?/AU

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DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N:y
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L7  ANSWER 1 OF 3  HCAPLUS  COPYRIGHT 2010 ACS on STN
ACCESSION NUMBER:  2004:515506  HCAPLUS
DOCUMENT NUMBER:   141:71453
TITLE:             Preparation of anthranilic acid amide derivatives as
                   neoplastic inhibitors
INVENTOR(S):       Bold, Guido; Furet, Pascal; Manley, Paul William
PATENT ASSIGNEE(S): Novartis Ag, Switz.; Novartis Pharma GmbH
SOURCE:            PCT Int. Appl., 81 pp.
                   CODEN: PIXXD2
DOCUMENT TYPE:      Patent
LANGUAGE:           English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:
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PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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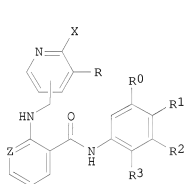
Updated Search

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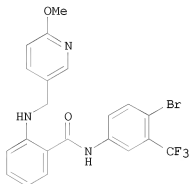
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RW: AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR				
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AU 2003294834	A1	20040630	AU 2003-294834	20031211
EP 1572686	A1	20050914	EP 2003-785795	20031211
EP 1572686	B1	20090415		
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BR 2003017292	A	20051108	BR 2003-17292	20031211
CN 1720244	A	20060111	CN 2003-80104845	20031211
CN 100427483	C	20081022		
JP 2006511518	T	20060406	JP 2004-558075	20031211
AT 428709	T	20090515	AT 2003-785795	20031211
PT 1572686	E	20090714	PT 2003-785795	20031211
ES 2324531	T3	20090810	ES 2003-785795	20031211
US 20060128684	A1	20060615	US 2005-538199	20050609
PRIORITY APPLN. INFO.:			GB 2002-29022	A 20021212
			WO 2003-EP14086	W 20031211

OTHER SOURCE(S): MARPAT 141:71453

GI



I



II

AB The title compds. I [wherein R and R0 = independently H, halo, (un)substituted alkyl, alkenyl, alkynyl, aryl, heteroaryl, etc.; R1 = H, halo, (un)substituted alkyl, alkenyl, alkynyl, alkoxy, OCF3, OCH2CF3, OCH2CH2CF3, or OCH2CH2CH2CF3; R2 = perfluoroalkyl; R3 = H or halo; X = OH, alkoxy, alkylthio, imino, alkylimino, halo, etc.; Z = N or CH] or salts, N-oxides, or tautomers thereof are prepared as neoplastic inhibitors for the treatment of human or animal body. For example, the compound II was prepared in a multi-step synthesis. Formulations containing I as an active ingredient were also described.

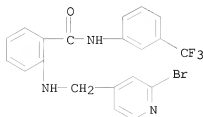
IT 657401-06-4P

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RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)
(drug candidate, reactant; preparation of anthranilic acid amide derivs. as neoplastic inhibitors)

RN 657401-06-4 HCAPLUS

CN Benzamide, 2-[(2-bromo-4-pyridinyl)methyl]amino]-N-[3-(trifluoromethyl)phenyl]- (CA INDEX NAME)



REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2010 ACS ON STN

ACCESSION NUMBER: 2004:120827 HCAPLUS

DOCUMENT NUMBER: 140:181330

TITLE: Preparation of anthranilamidopyridines as inhibitors of vascular endothelial growth factor receptor-2 and -3 (VEGFR-2 and -3).

INVENTOR(S): Huth, Andreas; Krueger, Martin; Zorn, Ludwig; Ince, Stuart; Thierauch, Karl-Heinz; Menrad, Andreas; Haberey, Martin; Hess-Stump, Holger

PATENT ASSIGNEE(S): Schering Aktiengesellschaft, Germany

SOURCE: PCI Int. Appl., 88 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004013102	A1	20040212	WO 2003-EP7964	20030722
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, BG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
DE 10235690	A1	20040219	DE 2002-10235690	20020731
DE 10328036	A1	20050105	DE 2003-10328036	20030619
CA 2493026	A1	20040212	CA 2003-2493026	20030722

Updated Search

AU 2003281855	A1	20040223	AU 2003-281855	20030722
BR 2003013122	A	20050705	BR 2003-13122	20030722
CN 1671666	A	20050921	CN 2003-818334	20030722
EP 1594841	A1	20051116	EP 2003-740470	20030722
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
JP 2005538112	T	20051215	JP 2004-525272	20030722
NZ 537291	A	20070223	NZ 2003-537291	20030722
US 20040147535	A1	20040729	US 2003-631018	20030731
US 7148357	B2	20061212		
US 20050054654	A1	20050310	US 2004-870491	20040618
US 7517894	B2	20090414		
MX 2004012948	A	20050912	MX 2004-12948	20041217
IN 2005DN00309	A	20070119	IN 2005-DN309	20050127
ZA 2005001642	A	20090930	ZA 2005-1642	20050224
NO 2005001035	A	20050429	NO 2005-1035	20050225
HR 2005000187	A2	20051031	HR 2005-187	20050225
US 20070015794	A1	20070118	US 2006-525091	20060922
US 7615565	B2	20091110		

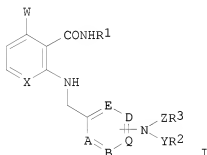
PRIORITY APPLN. INFO.:

DE 2002-10235690	A	20020731
DE 2003-10328036	A	20030619
US 2002-407970P	P	20020905
US 2003-483896P	P	20030702
WO 2003-EP7964	W	20030722
US 2003-631018	A3	20030731

ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT

OTHER SOURCE(S): MARPAT 140:181330

GI



AB Title compds. [I; X = CH, N; W = H, F; A, B, D, E, Q = N, C; ≤2 of A, B, D, E, Q = N; R1 = (substituted) aryl, heteroaryl; Y, Z = bond, CO, CS, SO2; R2, R3 = H, CONR9R10, SO2R6, COR11, NR9R10, (substituted) alkyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl; R2YNZAR3 = atoms to form a 3-8 membered (substituted) (unsatd.) ring; R6 = H, alkyl, haloalkyl, (substituted) aryl, heteroaryl, NR9R10; R9, R10 = H, alkyl, alkenyl, aryl, cycloalkyl, etc.; R11 = alkyl, alkoxy, hydroxyalkyl, hydroxyalkoxy, cycloalkyl, (substituted) Ph, pyridyl, biphenyl, naphthyl], were prepared Thus, 2-[(2-(2-bromopyridin-4-ylmethyl)amino)-N-(3-trifluoromethylphenyl)benzamide (preparation given) pyridine, and N,N-dimethylaminoethylamine were heated in a pressure vessel for 5 h at 200° to give 2-[(2-(2-dimethylaminoethylamino)pyridin-4-

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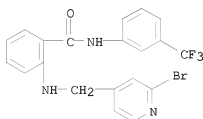
ylmethylamino]-N-(3-trifluoromethylphenyl)benzamide. I inhibited VEGFR-2 with IC50 = 8-65 nM. I can be used for treatment of tumor or metastasis growth, psoriasis, Kaposi's sarcoma, restenosis, endometriosis, Crohn's disease, Hodgkin's disease, leukemia, arthritis, hemangioma, angiofibroma, eye disease, renal diseases, transplant rejection, fibrotic diseases, mesangial cell proliferative diseases, atherosclerosis, injuries to nervous tissue and for inhibition of the reocclusion of vessels after balloon catheter treatment, in vessel prosthetics, or after the application of mech. devices to hold open vessels, as immunosuppressants, for scar-free wound healing, age spots and contact dermatitis.

IT 657401-06-4

RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation of anthranilamidopyridines as inhibitors of vascular endothelial growth factor receptor)

RN 657401-06-4 HCAPLUS

CN Benzamide, 2-[[2-bromo-4-pyridinyl)methyl]amino]-N-[3-(trifluoromethyl)phenyl]- (CA INDEX NAME)



OS.CITING REF COUNT: 18 THERE ARE 18 CAPLUS RECORDS THAT CITE THIS RECORD (20 CITINGS)
REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2010 ACS on STN

ACCESSION NUMBER: 2000:335388 HCAPLUS

DOCUMENT NUMBER: 132:347491

TITLE: Preparation of N-aryl(thio)anthranilic acid amides as VEGF receptor tyrosine kinase inhibitors
INVENTOR(S): Altmann, Karl-Heinz; Bold, Guido; Furet, Pascal; Manley, Paul William; Wood, Jeanette Marjorie; Ferrari, Stefano; Hofmann, Francesco; Mestan, Jurgen; Huth, Andreas; Kruger, Martin; Seidelmann, Dieter; Menrad, Andreas; Haberey, Martin; Thierauch, Karl-Heinz

PATENT ASSIGNEE(S): Novartis A.-G., Switz.; Novartis-Erfindungen Verwaltungsgesellschaft m.b.H.; Schering Aktiengesellschaft

SOURCE: PCT Int. Appl., 77 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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WO 2000027820      A1      20000518      WO 1999-EP8545      19991108
W:  AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU,
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    IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD,
    MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK,
    SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW
RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
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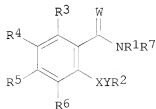
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EP 1129075      A1      20010905      EP 1999-971802    19991108
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HU 2001004188    A2      20020328      HU 2001-4188      19991108
HU 2001004188    A3      20020429
JP 2002529453    T      20020910      JP 2000-581000    19991108
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NZ 511339        A      20030725      NZ 1999-511339    19991108
CN 1152014        C      20040602      CN 1999-813108    19991108
RU 2286338        C2      20061027      RU 2001-114978    19991108
CZ 299829        B6      20081210      CZ 2001-1615     19991108
SK 287259        B6      20100407      SK 2001-628       19991108
NO 2001001894    A      20010704      NO 2001-1894      20010417
NO 328130        B1      20091214
ZA 2001003290    A      20030123      ZA 2001-3290      20010423
MX 2001004256    A      20030606      MX 2001-4256      20010427
US 20020019414   A1      20020214      US 2001-850434    20010507
US 6448277       B2      20020910
IN 2001CN00638   A      20050304      IN 2001-CN638     20010508
ZA 2001004673    A      20020909      ZA 2001-4673      20010607
US 20030064992   A1      20030403      US 2002-180289    20020626
US 6878720       B2      20050412
US 20040198782   A1      20041007      US 2004-828951    20040421
US 7002022       B2      20060221
US 20060074112   A1      20060406      US 2005-254897    20051020
GB 1998-24579    A      19981110
WO 1999-EP8545   W      19991108
US 2001-850434   A3      20010507
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US 2004-828951   A3      20040421

PRIORITY APPLN. INFO.:      MARPAT 132:347491

ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT
OTHER SOURCE(S):
GI

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AB Use of title compds. I; W = O, S; X = NR8; Y = CR9R10(CH2)n, SO2; R9, R10 = H, alkyl; n = 0-3; R1 = aryl; R2 = mono- or bicyclic heteroaryl with the exception that R2 cannot = 2-phthalimidyl, and when Y = SO2 cannot represent 2,1,3-benzothiadiazol-4-yl; R3-R6 = H, substituent; R7, R8 = H, alkyl; or a N-oxide or a pharmaceutically acceptable salt thereof, for the preparation of a pharmaceutical product for the treatment of a neoplastic disease which responds to an inhibition of the VEGF receptor tyrosine kinase activity is claimed. Thus, a mixture of 4-pyridinecarboxaldehyde and 2-amino-N-(4-trifluoromethylphenyl)benzamide (preparation given) in MeOH containing

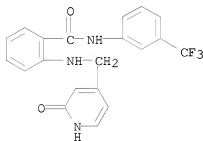
HOAc was treated with NaBH3CN followed by 16 h stirring to give 2-[(4-pyridyl)methyl]amino-N-[4-(trifluoromethyl)phenyl]benzamide. Tested I inhibited Flt-1 VEGF receptor tyrosine kinase with IC50 = 0.18-0.56 μ M.

IT 269391-01-7P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of N-aryl(thio)anthranilic acid amides as VEGF receptor tyrosine kinase inhibitors)

RN 269391-01-7 HCAPLUS

CN Benzamide, 2-[[[(1,2-dihydro-2-oxo-4-pyridinyl)methyl]amino]-N-[3-(trifluoromethyl)phenyl]- (CA INDEX NAME)



OS.CITING REF COUNT: 38 THERE ARE 38 CAPLUS RECORDS THAT CITE THIS RECORD (42 CITINGS)
REFERENCE COUNT: 14 THERE ARE 14 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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COST IN U.S. DOLLARS

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FULL ESTIMATED COST

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

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Updated Search

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DICTIONARY FILE UPDATES: 3 JUN 2010 HIGHEST RN 1226953-63-4

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TSCA INFORMATION NOW CURRENT THROUGH January 8, 2010.

Please note that search-term pricing does apply when conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

<http://www.cas.org/support/stngen/stdoc/properties.html>

=>

Uploading C:\Documents and Settings\brobinson1\My Documents\e-Red Folder\es6s6s6.str

L10 STRUCTURE UPLOADED

=> s l10

SAMPLE SEARCH INITIATED 16:31:29 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 1700 TO ITERATE

100.0% PROCESSED 1700 ITERATIONS

0 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 31527 TO 36473

PROJECTED ANSWERS: 0 TO 0

L11 0 SEA SSS SAM L10

=> s l10 full

THE ESTIMATED SEARCH COST FOR FILE 'REGISTRY' IS 191.05 U.S. DOLLARS

DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N or END:y

FULL SEARCH INITIATED 16:31:33 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 33645 TO ITERATE

100.0% PROCESSED 33645 ITERATIONS

2 ANSWERS

SEARCH TIME: 00.00.01

L12 2 SEA SSS FUL L10

=> d his

(FILE 'HOME' ENTERED AT 16:22:44 ON 05 JUN 2010)

Updated Search

stnvrkop

FILE 'REGISTRY' ENTERED AT 16:22:51 ON 05 JUN 2010

L1 STRUCTURE UPLOADED
L2 0 S L1
L3 0 S L1 FULL
L4 STRUCTURE UPLOADED
L5 0 S L4
L6 2 S L4 FULL

FILE 'HCAPLUS' ENTERED AT 16:27:33 ON 05 JUN 2010

L7 3 S L6
L8 0 S L7 AND LIGHTNER, J?/AU
L9 0 S L7 AND NG, H?/AU

FILE 'REGISTRY' ENTERED AT 16:29:11 ON 05 JUN 2010

L10 STRUCTURE UPLOADED
L11 0 S L10
L12 2 S L10 FULL

=> file hcaplus

COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	193.01	605.41
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	0.00	-2.55

FILE 'HCAPLUS' ENTERED AT 16:31:43 ON 05 JUN 2010

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FILE COVERS 1907 - 5 Jun 2010 VOL 152 ISS 24
FILE LAST UPDATED: 4 Jun 2010 (20100604/ED)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Apr 2010
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Apr 2010

HCAplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2010.

CAS Information Use Policies apply and are available at:

<http://www.cas.org/legal/infopolicy.html>

This file contains CAS Registry Numbers for easy and accurate

Updated Search

stnvrkop

substance identification.

=> s l12

L13 3 L12

=> d his

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L11 0 S L10

L12 2 S L10 FULL

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L13 3 S L12

=> s l13 not l7

L14 0 L13 NOT L7

Updated Search